

# STEDMAN'S

## Medical Dictionary

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immediately after sudden elevation of temperature; their function is to help diminish the harmful effects of high temperature.

**heterologous p.**, SYN foreign p.

**homologous p.'s**, p.'s having a very similar primary, secondary, and tertiary structure.

**immune p.**, SYN antibody.

**integral p.'s**, p.'s that cannot be easily separated from a biomembrane. SYN intrinsic p.'s.

**intrinsic p.'s**, SYN integral p.'s.

**iron-sulfur p.'s**, p.'s containing one or more iron atoms that are linked to sulfur bridges and/or sulfur of cysteinyl residues; e.g., certain p.'s in the electron transport pathway.

**p. kinases**, a class of enzymes that phosphorylates other p.'s; many of these kinases are responsive to other effectors (e.g., cAMP, cGMP, insulin, epidermal growth factor, calcium and calmodulin, calcium and phospholipids, etc.).

**M p.**, (1) SYN *Streptococcus M antigen*. SEE ALSO  $\beta$ -hemolytic *streptococci*, under *streptococcus*, *Streptococcus pneumoniae*. (2) SYN monoclonal immunoglobulin.

**macrophage inflammatory p.** (mak'rō-fāj in'flam-mā-to-rē), a member of the chemokine family that is chemotactic for certain lymphocyte subsets such as T cytotoxic cells.

**matrix Gla p.** (MGP), a calcium binding p.

**microtubule-associated p.'s (MAPs)**, p.'s that have a specific association with  $\alpha$ - and/or  $\beta$ -tubulin; e.g., 'tau, MAP1, MAP2; several have been found in the plaques observed in Alzheimer's disease.

**mild p. protein**, a complex prepared by the reaction of p. oxide with either gelatin or serum albumin. Black shiny crystals liberate p. and it was formerly widely used as a topical anti-infective on mucous membranes. Contains from 19 to 25% p., only a small fraction of which is ionizable. Can produce black or brown pigmentation due to deposition of reduced p. in the tissues. SYN argyrol, silvol.

**monoclonal p.**, SYN monoclonal immunoglobulin.

**monocyte chemoattractant p.-1 (MCP-1)** (mon'ō-sīt kē'mō-āk'trāk'tānt), secreted by endothelial cells of a blood vessel wall; it induces extravasation of monocytes.

**muscle p.'s**, p.'s present in muscle.

**myelin p. A1**, SYN encephalitogenic p.

**myeloblastic p.**, SEE human leukemia-associated *antigens*, under *antigen*.

**native p.**, the concept of a p. in its natural state, in the cell, unaltered by heat, chemicals, enzyme action, or the exigencies of extraction.

**neutrophil activating p. (NAP)**, SYN interleukin-8.

**non-heme iron p.**, any p. containing iron but not any heme iron; e.g., NADH dehydrogenase.

**nonspecific p.**, a p. substance that elicits a response not mediated by specific antigen-antibody reaction.

**odorant binding p.**, p.'s in nasal mucus that bind lipophilic odor-producing molecules and transfer them to the olfactory receptors. Similar p.'s may mediate taste.

**parathyroid hormonelike p. (PLP)**, a 140 amino acid p. secreted by some cancer cells; it causes hypercalcemia.

**pathological p.'s**, SEE paraprotein.

**peripheral p.'s**, p.'s that can be easily removed from a biomembrane (e.g., by altering the pH or the ionic strength). SYN extrinsic p.'s.

**phenylthiocarbamoyl p.**, formed by the reaction of phenylisothiocyanate with a terminal  $\alpha$ -amino group of a peptide or p. SEE ALSO phenylisothiocyanate, phenylthiohydantoin. SYN PhNCS p., PTC p.

**PhNCS p.**, SYN phenylthiocarbamoyl p.

**p. phosphatases**, a class of enzymes that catalyze the dephosphorylation of specific phosphorylated p.'s.

**placenta p.**, SYN human placental *lactogen*.

**plasma p.'s**, dissolved p.'s (more than 100) of blood plasma, mainly albumins and globulins (normally 6 to 8 g/100 ml); they hold fluid in blood vessels by osmosis and include antibodies and blood-clotting p.'s. SYN serum p.'s.

**prion p.**, small, infectious proteinaceous particle, of non-nucleic

acid composition because of its resistance to nucleases; the causative agent, either on a sporadic, genetic, or infectious basis, of six neurodegenerative diseases in animals, and four in humans; the latter include the spongiform encephalopathies of kuru, Creutzfeldt-Jakob disease, Gerstmann-Straussler-Scheinker syndrome and fatal familial insomnia. The gene encoding for the PrP is found on chromosome 20. SYN prion.

**protective p.**, SYN antibody.

**PTC p.**, SYN phenylthiocarbamoyl p.

**purified placental p.**, SYN human placental *lactogen*.

**receptor p.**, an intracellular p. (or p. fraction) that has a high specific affinity for binding a known stimulus to cellular activity, such as a steroid hormone or adenosine 3',5'-cyclic phosphate.

**retinol-binding p.**, a plasma p. that binds and transports retinol.

**S p.**, the major fragment produced from pancreatic ribonuclease by the limited action of subtilisin, which cleaves the ribonuclease between residues 20 and 21; the smaller fragment (residues 1-20) is S peptide.

**p. S**, a vitamin K-dependent antithrombotic p. that functions as a cofactor with activated p. C.

**serum p.'s**, SYN plasma p.'s.

**simple p.**, p. that yields only  $\alpha$ -amino acids or their derivatives by hydrolysis; e.g., albumins, globulins, glutelins, prolamines, albuminoids, histones, protamines. Cf. conjugated p.

**stimulatory p. 1 (SP1)**, an RNA polymerase II transcription factor in vertebrates; binds to DNA in regions rich in G and C residues; a general promoter-binding factor necessary for the activation of many genes.

**strong silver p.**, SEE strong *silver protein*.

**structure p.'s**, p.'s whose role is for structure and support in tissue and within the cell; e.g., the collagens.

**Tamm-Horsfall p.**, SEE Tamm-Horsfall *mucoprotein*.

**thyroxine-binding p. (TBP)**, (1) SYN thyroxine-binding *globulin*. (2) SYN thyroxine-binding *prealbumin*.

**unwinding p.'s**, enzymes that uncoil the DNA allowing recombination events to occur.

**vitamin D-binding p. (DBP)**, a plasma p. that binds vitamin D.

**whew p.**, the soluble p. contained in the whey of milk clotted by rennin; e.g., lactoglobulin,  $\alpha$ -lactalbumin, lactoferrin.

**Z-p.**, a fatty acid-binding protein that participates in the intracellular movement of fatty acids. SYN fatty acid binding p.

**pro-tein-a-ceous** (prō'tē-nā-shūs, prō'tē-i-nā-shūs). Resembling a protein; possessing, to some degree, the physicochemical properties characteristic of proteins.

**pro-tein-ase**. SYN endopeptidase.

**pro-tein hy-drol-y-sate**. A sterile solution of amino acids and soft chain peptides prepared from a suitable protein by acid or enzymatic hydrolysis; used intravenously for the maintenance of positive nitrogen balance in severe illness, and after surgery involving the alimentary tract; or used orally in the diets of infants allergic to milk or as a supplement when high protein intake from ordinary foods cannot be accomplished.

**pro-tein-o-gen-ic** (prō'ten-ō-jen'ik). SYN proteogenic.

**pro-tein-oids** (prō'tēn-oydz; prō'tē-in-oydz). Artificially synthesized heteropoly(amino acids).

**pro-tein-o-sis** (pro-tē-nō'sis, prō'tē-i-nō'sis). A state characterized by disordered protein formation and distribution, particularly as manifested by the deposition of abnormal proteins in tissues: [protein + G. -osis, condition]

**lipoid p.** [MIM\*247100], a disturbance of lipid metabolism in which there are deposits of a protein-lipid complex on the oral tongue and sublingual and faucial areas, and translucent keratotic papillomatous eyelid lesions; autosomal recessive inheritance, frequently with intracranial calcifications. SYN hyalinosis cutis et mucosae, lipoidosis cutis et mucosae, Urbach-Wiethe disease.

**pulmonary alveolar p.**, a chronic progressive lung disease of adults, characterized by alveolar accumulation of granular proteinaceous material that is PAS-positive and lipid rich, with little inflammatory cellular exudate; the cause is unknown.

**pro-tein-u-ria** (prō-tē-nū'rē-ā, prō'tē-i-nū'rē-ā). 1. Presence of urinary protein in concentrations greater than 0.3 g in a 24-hour urine collection or in concentrations greater than 1 g/l (1+ to 2+